FILE 'REGISTRY' ENTERED AT 15:23:36 ON 03 MAR 2004 L1 109 S AGVDNRECI | AQIFNKPYW/SQSP

FILE 'HCAPLUS' ENTERED AT 15:24:49 ON 03 MAR 2004 L2 30 S L1

L4 22 S L2(L) (HPV OR PAPILLOM? OR WART VIRUS)

L4 ANSWER 1 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 11 Jan 2004

ACCESSION NUMBER: 2004:20437 HCAPLUS

DOCUMENT NUMBER: 140:87675

TITLE: Antisense oligonucleotides, ribozymes and

DNAzymes targeting human papillomavirus genes E6 & E7 for treatment of HPV infections associated

with cervical cancer

INVENTOR(S): Clawson, Gary A.; Pan, Wei-Hua; Christensen,

Neil; Thiboutot, Diane

PATENT ASSIGNEE(S): The Penn State Research Foundation, USA

SOURCE: PCT Int. Appl., 65 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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PATENT NO.				KIND DATE		APPLICATION NO. DATE										
WO 2	0 2004002416			A2 20040108			WO 2003-US2034						40 20030626			
	W:	ΑE,	AG,	AL,	AM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,
		CN,	co,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KP,	KR,	ΚZ,
		LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,
		NO,	NZ,	OM,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	ТJ,
		TM,	TN,	TR,	TT,	TZ,	UA,	ŪG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW,
		AM,	AZ,	BY,	KG,	ΚZ,	MD,	RU,	ΤJ,	TM						
	RW:	GH,	GM,	ΚE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	ΑT,	BE,
		BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	IT,
		LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,
		GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG.					
PRIORITY	APPI	LN.	INFO	. :				1	US 20	002-	3917	95P	P	2002	0626	
								1	US 20	002-	4179	97P	P	2002	1014	
								1	US 20	003-	4490	66P	P	2003	0221	

The invention provides antisense oligonucleotides (ASO), ribozymes (Rz) and DNAzymes (Dz) targeting human papillomavirus (HPV) genes E6 & E7 for treatment of HPV infections associated with cervical cancer. The ASO, Rz, or Dz are provided in a topical drug, in conjunction with keratolytic agents (salicylic acid), and applied to the HPV infected cells (cervical, skin or epithelium). The ASO, Rz or Dz target HPV genes E6/E7 mRNA for cleavage, resulting in a decreased replication rate and thus decreased number of cells infected. The invention can be applied towards treating HPV infections (e.g., HPV infections of cutaneous and mucosal epithelial cells) and HPV-associated conditions (e.g., cervical dysplasia, HPV-associated cervical carcinomas, oral mucosal papilloma cancers, laryngeal papilloma cancers) in humans.

IT 624440-70-6

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RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)

(amino acid sequence; antisense oligonucleotides, ribozymes and DNAzymes targeting HPV genes E6 & E7 for treatment of HPV infections associated with cervical cancer)

L4 ANSWER 2 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 28 Nov 2003

ACCESSION NUMBER: 2003:931391 HCAPLUS

DOCUMENT NUMBER: 140:1544

TITLE: Methods for preparing chimeric human

papillomavirus 16 L1 virus like particles and

uses as vaccines

INVENTOR(S):

Varsani, Arvind Devshi; Rybicki, Edward Peter

APPLICATION NO.

WO 2003-IB1912

20030519

PATENT ASSIGNEE(S): University of Cape Town, S. Afr.

SOURCE:

PCT Int. Appl., 32 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

DATE

20031127

KIND

A2

627561-79-9P 627561-80-2P 627561-81-3P

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.

WO 2003097673

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W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD,
             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ,
             LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
             NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ,
             TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW,
             AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE,
             BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT,
             LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM,
             GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                                        ZA 2002-3957
                                                         A 20020517
PRIORITY APPLN. INFO.:
     The invention describes a method for producing a chimeric human
    papillomavirus (HPV) L1 polypeptide containing a heterologous peptide,
     and in particular, a HPV L2 peptide. The method comprises the steps
     of introducing a DNA sequence coding for the heterologous peptide
     into a DNA sequence coding for the L1 polypeptide; introducing the
    DNA sequence including the sequences for the L1 polypeptide and
    heterologous peptide into a host cell in which the DNA sequence can
    be expressed; causing expression of the DNA sequence; and recovering
     the resulting chimeric L1 polypeptide which includes the
    heterologous peptide. Typically, the nucleotides encoding the
    heterologous peptide replace the nucleotides of the L1 polypeptide
     at the point of insertion. The invention also describes a vector
     for use in the method, a host cell containing the vector, and a vaccine
     including the chimeric HPV L1 polypeptide produced according to the
    method.
IT
     627561-74-4P 627561-77-7P 627561-78-8P
```

Searcher: Shears 571-272-2528

RL: BPN (Biosynthetic preparation); BSU (Biological study,

unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (amino acid sequence; methods for preparing chimeric human papillomavirus 16 L1 virus like particles and uses as vaccines)

ANSWER 3 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN T.4

Entered STN: 14 Nov 2003

ACCESSION NUMBER: 2003:892896 HCAPLUS

139:379997 DOCUMENT NUMBER:

Production of transgenic plant expressing TITLE:

papillomavirus L1 capsid protein and use as

APPLICATION NO. DATE

papillomavirus vaccines

Rose, Robert C.; Mason, Hugh S.; Warzecha, INVENTOR(S):

Heribert

University of Rochester, USA; Boyce Thompson PATENT ASSIGNEE(S):

Institute for Plant Research, Inc.

SOURCE: PCT Int. Appl., 74 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

LANGUAGE:

i

Patent English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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PATENT NO.
                         KIND DATE
                                                   WO 2003-US13757 20030502
      WO 2003093437 A2
                                  20031113
           W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,
                CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD,
               GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM,
                ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
           RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT,
                LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM,
                GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                                                  US 2002-377467P P 20020502
PRIORITY APPLN. INFO.:
      The present invention relates to a method of producing
      papillomavirus virus-like particles or capsomeres. This method
      involves providing a transgenic plant or plant seed transformed with
      a nucleic acid mol. comprising a papillomavirus L1 capsid protein
      coding sequence and growing the transgenic plant or a transgenic
      plant grown from the transgenic plant seed under conditions
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IT 622878-55-1

RL: PRP (Properties)

(unclaimed protein sequence; production of transgenic plant expressing papillomavirus L1 capsid protein and use as papillomavirus vaccines)

can be administered to a subject under conditions effective to immunize the subject against disease resulting from infection by a papillomavirus. DNA constructs, expression vectors, host cells,

plants, and plant seeds are also disclosed.

571-272-2528 Searcher : Shears

effective to produce virus-like particles containing the papillomavirus L1 capsid protein. The plant or a component part or a fruit thereof

i

INVENTOR(S):

ANSWER 4 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN Entered STN: 22 Aug 2003 ACCESSION NUMBER: 2003:656897 HCAPLUS 139:193297 DOCUMENT NUMBER: Complexes of human and human papillomavirus TITLE: proteins and their use in drug screening and diagnosis Jackson, Amanda; Ooi, Chean Eng; Lewin, David INVENTOR(S): A.; Cuthill, Scott Curagen Corporation, USA; Hoffmann-La Roche Inc. PATENT ASSIGNEE(S): SOURCE: PCT Int. Appl., 156 pp. CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: KIND DATE APPLICATION NO. DATE PATENT NO. WO 2003-US4594 20030214 A2 20030821 WO 2003068940 **A**3 20031127 WO 2003068940 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG US 2002-356911P P 20020214 PRIORITY APPLN. INFO.: Complexes of human papillomavirus (HPV) proteins E1-E7, L1, and L2 with human proteins are disclosed. These complexes may be used to screen for agents which disrupt the complexes. These agents may be used for treatment of HPV infections. A method of detecting these complexes may be used in screening for pre-cancerous cervical lesions and for classifying HPV infections. Thus, yeast two-hybrid assays were used to identify interactions of HPV-la, HPV-11, and HPV-16 proteins with human proteins. IT 583085-22-7 RL: PRP (Properties) (unclaimed protein sequence; complexes of human and human papillomavirus proteins and their use in drug screening and diagnosis) ANSWER 5 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN L4Entered STN: 22 Aug 2003 2003:656523 HCAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 139:196258 Manufacture of papillomavirus-like particles for TITLE: vaccine use in insect cells using codon-optimized synthetic genes

Searcher: Shears 571-272-2528

Robinson, Robin A.; Cioce, Vittoria

PATENT ASSIGNEE(S):

Novavax, Inc., USA

SOURCE:

PCT Int. Appl., 126 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent English

LANGUAGE:

COLINIT. 4

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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APPLICATION NO.
                                                            DATE
     PATENT NO.
                      KIND
                            DATE
                                           ______
                                                            20030214
     WO 2003068163
                            20030821
                                          WO 2003-US4473
                      A2
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD,
             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ,
             LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
             NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ,
             TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW,
             AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE,
             BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT,
             LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA,
             GN, GQ, GW, ML, MR, NE, SN, TD, TG
                            20031211
                                           US 2003-367095
                                                            20030214
     US 2003228696
                      A1
PRIORITY APPLN. INFO.:
                                        US 2002-356113P P
                                                            20020214
                                        US 2002-356118P P
                                                            20020214
                                        US 2002-356119P P
                                                            20020214
                                        US 2002-356123P P
                                                            20020214
                                        US 2002-356126P P
                                                            20020214
                                        US 2002-356133P P
                                                            20020214
                                        US 2002-356135P
                                                        Ρ
                                                            20020214
                                        US 2002-356150P
                                                        Ρ
                                                            20020214
                                        US 2002-356151P P
                                                            20020214
                                        US 2002-356152P
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                                                            20020214
                                        US 2002-356154P
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                                                            20020214
                                                        P
                                        US 2002-356156P
                                                            20020214
                                        US 2002-356157P
                                                        Ρ
                                                            20020214
                                                        Ρ
                                        US 2002-356161P
                                                            20020214
                                        US 2002-356162P P
                                                            20020214
     Virus-like particles that exhibit conformational antigenic epitopes
AB
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capable of eliciting neutralizing antibodies to human papillomaviruses are manufactured in insect cells using synthetic genes with codon usage optimized for efficient expression of these genes in animal cells. Pharmaceutical compns., vaccines, and diagnostic test kits containing the chimeric virus-particles are also provided. The proteins that form the virus-like particles are manufactured by expression of the genes in a derivative of the Sf9 cell line (Sf9S) using a baculovirus expression vector. Manufacture and chromatog. purification of human papillomavirus 16 virus-like particles is demonstrated. Vaccines containing the virus-like particles were well-tolerated by human volunteers and vaccines using Mf-59 adjuvant showed a prolonged immunity to the virus.

IT 582340-39-4

RL: PRP (Properties)

(unclaimed protein sequence; manufacture of **papillomavirus** -like particles for vaccine use in insect cells using codon-optimized synthetic genes)

L4 ANSWER 6 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 16 Apr 2003

ACCESSION NUMBER: 2003:291123 HCAPLUS

DOCUMENT NUMBER: 138:383747

TITLE: Human papillomavirus type 16 L1 capsomeres

induce L1-specific cytotoxic T lymphocytes and

tumor regression in C57BL/6 mice

AUTHOR(S): Ohlschlager, Peter; Osen, Wolfram; Dell,

Kerstin; Faath, Stefan; Garcea, Robert L.;
Jochmus, Ingrid; Muller, Martin; Pawlita,
Michael; Schafer, Klaus; Sehr, Peter; Staib,

Caroline; Sutter, Gerd; Gissmann, Lutz

CORPORATE SOURCE: Angewandte Tumorvirologie, Deutsches

Krebsforschungszentrum, Heidelberg, D-69120,

Germany

SOURCE: Journal of Virology (2003), 77(8), 4635-4645

CODEN: JOVIAM; ISSN: 0022-538X

PUBLISHER: American Society for Microbiology

DOCUMENT TYPE: Journal LANGUAGE: English

We analyzed capsomeres of human papillomavirus type 16 (HPV16) consisting of the L1 major structural protein for their ability to trigger a cytotoxic T-cell (CTL) response. To this end, we immunized C57BL/6 mice and used the L1165-173 peptide for ex vivo restimulation of splenocytes prior to anal. (51Cr release assay and enzyme-linked immunospot assay [ELISPOT]). This peptide was identified in this study as a Db-restricted naturally processed CTL epitope by HPV16 L1 sequence anal., major histocompatibility complex class I binding, and 51Cr release assays following immunization of C57BL/6 mice with HPV16 L1 virus-like particles (VLPs). HPV16 L1 capsomeres were obtained by purification of HPV16 L1 lacking 10 N-terminal amino acids after expression in Escherichia coli as a glutathione S-transferase fusion protein (GST-HPV16 $L1\Delta N10$). Sedimentation anal. revealed that the majority of the purified protein consisted of pentameric capsomeres, and assembled particles were not observed in minor contaminating higher-mol.-weight material. S.c. (s.c.) as well as intranasal immunization of C57BL/6 mice with HPV16 L1 capsomeres triggered an L1-specific CTL response in a dose-dependent manner as measured by ELISPOT and 51Cr release assay. Significant reduction of contaminating bacterial endotoxin (lipopolysaccharide) from the capsomere preparation did not diminish the immunogenicity. Antibody responses (serum and vaginal) were less robust under the exptl. conditions employed. In addition, s.c. vaccination with HPV16 L1 capsomeres induced regression of established tumors expressing L1 determinants (C3 tumor cells). data demonstrate that capsomeres are potent inducers of CTL responses similar to completely assembled T=7 VLPs. This result is of potential relevance for the development of (combined prophylactic and therapeutic) HPV-specific vaccines, since capsomeres can be produced easily and also can be modified to incorporate heterologous sequences such as early HPV proteins.

IT 308800-09-1

RL: BSU (Biological study, unclassified); BIOL (Biological study) (human papillomavirus type 16 L1 capsomeres induce L1-specific cytotoxic T lymphocytes and tumor regression in

C57BL/6 mice)

REFERENCE COUNT: 51 THERE ARE 51 CITED REFERENCES AVAILABLE

FOR THIS RECORD. ALL CITATIONS AVAILABLE

IN THE RE FORMAT

L4 ANSWER 7 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 15 Nov 2002

ACCESSION NUMBER: 2002:869424 HCAPLUS

DOCUMENT NUMBER: 137:364445

TITLE: DNA sequences encoding human papillomavirus 16

L1 proteins capable of efficiently forming

virus-like particles

INVENTOR(S): Durst, Matthias; Gissmann, Latz

PATENT ASSIGNEE(S): Germany

SOURCE: U.S. Pat. Appl. Publ., 17 pp., Cont. of U.S.

Ser. No. 884,168, abandoned.

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

LANGUAGE: Eng FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

US 2002168372 A1 20021114 US 1998-162904 19980929

PRIORITY APPLN. INFO.: US 1993-92528 B1 19930716

US 1996-641570 B1 19960501

US 1997-884168 B1 19970627

- The present invention provides novel DNA sequences encoding human papillomavirus L1 proteins capable of efficiently forming virus-like particles (VLP). In particular, the present invention relates to a DNA sequence encoding an HPV16 L1 protein. Furthermore, the present invention relates to expression plasmids containing said DNA, to host cells transformed by said expression plasmids, to methods for the production of said L1 protein, to the VLP formed by said L1 protein, to antibodies reacting with said protein and said VLP, to diagnostic and pharmaceutical compns. and methods and to a vaccine comprising said VLP.
- 1T 475437-53-7P, Protein L1 (human papillomavirus 16)
 475437-55-9P, Protein L1 (human papillomavirus 16)
 RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(amino acid sequence; DNA sequences encoding human papillomavirus 16 L1 proteins capable of efficiently forming virus-like particles)

L4 ANSWER 8 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 07 Jun 2002

ACCESSION NUMBER: 2002:429086 HCAPLUS

DOCUMENT NUMBER: 137:19374

TITLE: T-cell epitopes of papillomavirus L1 and E7

proteins and their use in diagnosis and therapy

of infection

INVENTOR(S): Nieland, John; Kaufmann, Andreas

PATENT ASSIGNEE(S): Medigene Aktiengesellschaft, Germany; Kather,

Angela; Schinz, Manuela PCT Int. Appl., 126 pp.

SOURCE: PCT Int. Appl CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PAT	TENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO	2002044384	A2	20020606	WO 2001-EP14037	20011130
WO	2002044384	A 3	20031016		
	W: AU, CA,	JP, US			
	RW: AT, BE,	CH, CY	DE, DK, ES	, FI, FR, GB, GR, IE,	, IT, LU, MC,
	NL, PT,	SE, TR			
DΕ	10059631	A1	20020718	DE 2000-10059631	
ΑU	2002020744	A 5	20020611	AU 2002-20744	
EP		•	20031217	EP 2001-998642	
	R: AT, BE,	CH, DE	DK, ES, FR	, GB, GR, IT, LI, LU,	, NL, SE, MC,
	PT, IE,	FI, CY	TR		
~		_		DE 2000 10050621 X	20001201

PRIORITY APPLN. INFO.:

DE 2000-10059631 A 20001201 WO 2001-EP14037 W 20011130

AB T cell epitopes of papillomavirus L1 and E7 proteins that may be useful in the diagnosis and treatment of viral infection are identified and characterized. Identification of epitopes using chimeric baculovirus-like particles is demonstrated.

IT 434341-62-5 434341-76-1

RL: PRP (Properties)

(unclaimed sequence; t-cell epitopes of papillomavirus L1 and E7 proteins and their use in diagnosis and therapy of infection)

L4 ANSWER 9 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 27 Sep 2001

ACCESSION NUMBER: 2001:705899 HCAPLUS

DOCUMENT NUMBER: 136:257935

TITLE: Enhancement of capsid gene expression: preparing

the human papillomavirus type 16 major

structural gene L1 for DNA vaccination purposes

AUTHOR(S): Leder, Christoph; Kleinschmidt, Jurgen A.;

Wiethe, Carsten; Muller, Martin

CORPORATE SOURCE: Forschungsschwerpunkt fur Angewandte

Tumorvirologie, Deutsches Krebsforschungszentrum

Heidelberg, Heidelberg, 69120, Germany

SOURCE: Journal of Virology (2001), 75(19), 9201-9209

CODEN: JOVIAM; ISSN: 0022-538X

PUBLISHER: American Society for Microbiology

DOCUMENT TYPE: Journal LANGUAGE: English

AB Expression of the structural proteins L1 and L2 of the human papillomaviruses (HPV) is tightly regulated. As a consequence, attempts to express these prime-candidate genes for prophylactic vaccination against papillomavirus-associated diseases in mammalian cells by means of simple DNA transfections result in insufficient production of the viral antigens. Similarly, in vivo DNA vaccination

using HPV L1 or L2 expression constructs produces only weak immune responses. In this study we demonstrate that transient expression of the HPV type 16 L1 and L2 proteins can be highly improved by changing the RNA coding sequence, resulting in the accumulation of significant amts. of virus-like particles in the nuclei of transfected cells. Data presented indicate that, in the case of L1, adaptation for codon usage accounts for the vast majority of the improvement in protein expression, whereas translation-independent posttranscriptional events contribute only to a minor degree. Finally, the adapted L1 genes demonstrate strongly increased immunogenicity in vivo compared to that of unmodified L1 genes.

IT 405194-67-4P

RL: BPN (Biosynthetic preparation); PRP (Properties); BIOL (Biological study); PREP (Preparation)

(amino acid sequence; enhancement of capsid gene expression, preparing the human papillomavirus type 16 major structural gene L1 for DNA vaccination purposes)

REFERENCE COUNT:

THERE ARE 44 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 10 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

44

ED Entered STN: 15 Jun 2001

ACCESSION NUMBER: 2001:434895 HCAPLUS

DOCUMENT NUMBER: 135:45179

TITLE: Inducing cellular immune responses to human

papillomavirus using peptide and nucleic acid

compositions

INVENTOR(S): Sette, Alessandro; Sidney, John; Southwood,

Scott; Chesnut, Robert; Celis, Esteban; Grey,

Howard M.

PATENT ASSIGNEE(S): Epimmune Inc., USA

SOURCE: PCT Int. Appl., 756 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA'	PENT	NO.		KI:	ND .	DATE			A.	PPLI	CATI	ON N	ο.	DATE		
WO 2001041799			99	A1 20010614					WO 2000-US33549 20001211							
	W:													BZ,		
														GD,		
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	LK,
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		PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TR,	TT,	ΤZ,
		UA,	ŪĠ,	US,	·UZ,	VN,	YU,	ZA,	ZW,	AM,	ΑZ,	BY,	KG,	KZ,	MD,	RU,
		ТJ,	TM													
	RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	ΤZ,	ŪG,	ZW,	AT,	BE,	CH,
		CY,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,
		TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GW,	ML,	MR,	NE,	SN,	TD,
		TG														
ΕP	1246															
	R:												LU,	NL,	SE,	MC,
		PT,	ΙE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR				

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US 1999-172705P P 19991210
PRIORITY APPLN. INFO.:
                                       US 2000-641528 A 20000815
                                       WO 2000-US33549 W 20001211
     This invention uses our knowledge of the mechanisms by which antigen
AB
     is recognized by T cells to identify and prepare human papillomavirus
     (HPV) epitopes, and to develop epitope-based vaccines directed
     towards HPV. More specifically, this application communicates our
     discovery of pharmaceutical compns. and methods of use in the
     prevention and treatment of HPV infection. The disclosed human
    papillomavirus protein epitopes include HLA-A1, HLA-A2, HLA-A3,
     HLA-A24, HLA-B7, HLA-B27, HLA-B58, HLA-B62 and HLA-DR supermotifs;
     and HLA-A1, HLA-A2, HLA-A3, HLA-A24, HLA-A11, HLA-DR3a and HLA-DR3b
    motifs. These supermotifs and motifs are derived from E1, E2, E5,
     E6, E7, L1 and L2 proteins of HPV16, HPV18, HPV31, HPV33, HPV45,
    HPV56, HPV6A, HPV6B, and HPV11.
     243135-53-7 344806-30-0, Protein L1 (human
    papillomavirus 31)
    RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (amino acid sequence; epitope-based vaccines for inducing
        cellular immune responses to human papillomavirus)
                              THERE ARE 8 CITED REFERENCES AVAILABLE FOR
                         8
REFERENCE COUNT:
                               THIS RECORD. ALL CITATIONS AVAILABLE IN
                               THE RE FORMAT
    ANSWER 11 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN
    Entered STN: 12 Dec 2000
                         2000:865093 HCAPLUS
ACCESSION NUMBER:
                         134:16537
DOCUMENT NUMBER:
                         Cytotoxic T-cell epitopes of the Papillomavirus
TITLE:
                         L1-Protein and their use in diagnostics and
                         therapy
                         Schaefer, Klaus; Faath, Stefan; Jochmus, Ingrid;
INVENTOR(S):
                         Nieland, John; Osen, Wolfram
                         Medigene A.-G., Germany; Deutsches
PATENT ASSIGNEE(S):
                         Krebsforschungszentrum (DKFZ)
                         Ger. Offen., 26 pp.
SOURCE:
                         CODEN: GWXXBX
DOCUMENT TYPE:
                         Patent
                         German
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
                                          APPLICATION NO. DATE
                    KIND DATE
     PATENT NO.
                    ____
                           _____
                            20001207
                                          DE 1999-19925235 19990601
     DE 19925235
                      A1
                                          WO 2000-EP5005 20000531
                            20001207
     WO 2000073464
                      A1
         W: AU, CA, JP, US
         RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,
             NL, PT, SE
                            20020306
                                          EP 2000-936846
                                                            20000531
     EP 1183367
                      A1
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
             PT, IE, FI
                            20030128
                                           JP 2001-500776
                                                            20000531
     JP 2003503018
                       Т2
PRIORITY APPLN. INFO.:
                                        DE 1999-19925235 A 19990601
                                        WO 2000-EP5005 W 20000531
```

The available invention concerns Papillomavirus T-cell epitopes with AΒ an amino acid sequence AQIFNKPYW, AGVDNRECI, and/or a functionally active variant thereof, as well as their use in diagnostics and therapy. 308800-07-9, AQIFNKPYW peptide+ 308800-09-1, IT AGVDNRECI peptide+ RL: BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses) (cytotoxic T-cell epitopes of Papillomavirus L1 protein and use in diagnostics and therapy) THERE ARE 3 CITED REFERENCES AVAILABLE FOR REFERENCE COUNT: 3 THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 12 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN L4Entered STN: 22 Sep 2000 2000:666563 HCAPLUS ACCESSION NUMBER: 133:256735 DOCUMENT NUMBER: Human papillomavirus L1 proteins for use in TITLE: vaccines, diagnostic reagents, and tools for studying surface receptor interactions Harrison, Stephen; Chen, Xiaojiang INVENTOR(S): The President & Fellows of Harvard College, USA PATENT ASSIGNEE(S): PCT Int. Appl., 31 pp. SOURCE: CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: KIND DATE APPLICATION NO. DATE PATENT NO. ____ _____ _____ 20000308 20000921 WO 2000-US6017 WO 2000054730 A2 20011115 WO 2000054730 A3 AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG 20030422 20000308 US 2000-520822 US 6551597 B1 US 2002-301260 20021121 20030626 US 2003118609 Α1 US 1999-125208P P 19990318 PRIORITY APPLN. INFO.:

AB Large quantities of soluble multimers of human papillomavirus L1 proteins can be produced in bacterial expression systems and used as therapeutic and diagnostic tools. L1 multimers can be used in immunogenic vaccine compns., as diagnostic reagents, and as tools for mapping cell surface receptor interactions.

IT 295372-67-7P, Protein L1 (human papillomavirus 16)

US 1999-148544P P 19990812

A3 20000308

US 2000-520822

RL: BAC (Biological activity or effector, except adverse); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PEP (Physical, engineering or chemical process); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses)

(amino acid sequence; human papillomavirus L1 proteins for use in vaccines, diagnostic reagents, and tools for studying surface receptor interactions)

L4 ANSWER 13 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 04 Jun 2000

ACCESSION NUMBER: 2000:368423 HCAPLUS

DOCUMENT NUMBER: 133:22397

TITLE: Chimeric biotin-binding papillomavirus protein

for delivery of biotinylated compounds to cells

Mueller, Martin; Kast, Wijbe M.; Nieland, John

D.; Velders, Markwin P.

PATENT ASSIGNEE(S): Loyola University of Chicago, USA

SOURCE: PCT Int. Appl., 27 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

INVENTOR(S):

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APPLICATION NO. DATE
                              KIND DATE
       PATENT NO.
                                       _____
                                                         WO 1999-US27555 19991122
                                       20000602
                             A1
      WO 2000031128
            W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA,
                  ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
            RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
                  DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF,
                  BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                                            US 1999-413611
                                                                                     19991006
                                B1 20020430
       US 6380364
                                                        US 1998-109510P P 19981123
PRIORITY APPLN. INFO.:
                                                        US 1999-413611
                                                                               A 19991006
```

AB The present invention provides a chimeric protein including a first domain which includes at least a portion of a papillomavirus L1 or L2 protein and a second domain which includes a biotin-binding polypeptide. The invention also provides papillomaviruses, capsomeres, and VLPs (virus-like particles), including such chimeric proteins and a method for delivering biotinylated substances to cells using such reagents.

IT 272100-89-7 272100-91-1

RL: BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)

(amino acid sequence; chimeric biotin-binding papillomavirus protein for delivery of biotinylated compds. to cells)

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 14 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 01 Oct 1999

ACCESSION NUMBER: 1999:626218 HCAPLUS

DOCUMENT NUMBER: 131:262610

TITLE: Formulation having a papilloma virus-specific

protein

INVENTOR(S): Burger, Alexander; Gabelsberger, Josef PATENT ASSIGNEE(S): Medigene Aktiengesellschaft, Germany

SOURCE: PCT Int. Appl., 26 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	KI	KIND DATE				7	APPL	ο.	DATE										
	WO	9948	 917			2	1999	0930		V	70 19	999-E	P199	9	1999	0324			
	WO	9948917			A	3	19991209												
			AU,																
		RW:	AT,	BE,	CH,	CY,	DE,	DK,	ES,	FI,	FR,	, GB,	GR,	IE,	IT,	LU,	MC,		
			NL,	PT,	SE														
	DE	1981	2940		A	_	1999			_		998-1			1998				
	DE	2982	4556				2001			_		998-2	-						
	CA	2323	526				1999					999-2							
	AU	9935	989			_	1999			-		999-3							
	ΕP	1066			A.	_	2001					999-9							
		R:	AT,	ΒE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	, IT,	LI,	LU,	NL,	SE,	MC,		
				ΙE,										_					
	JP	2002	50762	25	T	2	2002	0312				000-5			1999				
PRIC	RIT	APP	LN.	INFO	.:							-1981			1998				
										WO :	L999-	-EP19	99	W	1999	J324			

AB Papilloma virus-specific early and late proteins are soluble in formulations containing 0.3-.apprx.4M salt with pH 7.3-7.45. Formulations containing these proteins, as well as deletion mutants and chimeric proteins which form viruslike particles, are useful for therapeutic and diagnostic purposes. Thus, a human papilloma virus 16 L1E7 fusion protein gene was constructed by recombinant DNA technol., and expressed as viruslike particles in Trichoplusia ni cells using a baculovirus vector.

IT 244773-43-1

RL: BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)

(formulation having a papilloma virus-specific protein)

L4 ANSWER 15 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 27 Jul 1999

ACCESSION NUMBER: 1999:457683 HCAPLUS

DOCUMENT NUMBER: 131:209883

TITLE: Nucleotide sequences and further

characterization of human papillomavirus DNA

present in the CaSki, SiHa and HeLa cervical

carcinoma cell lines

AUTHOR(S):

Meissner, John D.

CORPORATE SOURCE:

Human Papillomavirus Section, National Center for Infectious Diseases, Centers for Disease Control and Prevention, Public Health Service, US Department of Health and Human Services,

Atlanta, GA, 30333, USA

SOURCE:

Journal of General Virology (1999), 80(7),

1725-1733

CODEN: JGVIAY; ISSN: 0022-1317 Society for General Microbiology

DOCUMENT TYPE:

Journal

LANGUAGE:

PUBLISHER:

English

The complete nucleotide sequences of the human papillomavirus type 16 (HPV-16) variants present in the CaSki and SiHa cervical carcinoma cell lines and the primary subgenomic HPV-18 variant present in the HeLa cervical carcinoma cell line were determined using overlapping bulk PCR products as templates. PCR-based methods were also used to characterize five previously unreported CaSki HPV-16 genomic disruptions and the 5' cellular-viral junction common to all HeLa HPV-18 subgenomic structures.

243135-53-7 IT

RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)

(amino acid sequence; nucleotide sequences and further characterization of human papillomavirus DNA present in CaSki, SiHa and HeLa cervical carcinoma cell lines)

REFERENCE COUNT:

THERE ARE 60 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE

IN THE RE FORMAT

ANSWER 16 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN L4

60

Entered STN: 02 Feb 1998

ACCESSION NUMBER: 1998:61423 HCAPLUS

DOCUMENT NUMBER:

128:213916

TITLE:

The genomes of three of four novel HPV types,

defined by differences of their L1 genes, show high conservation of the E7 gene and the URR Delius, Hajo; Saegling, Bettina; Bergmann, Krister; Shamanin, Vladimir; De Villiers,

Ethel-Michele

CORPORATE SOURCE:

Division for Tumorvirus Characterization, Research Program Applied Tumorvirology, Deutsches Krebsforschungszentrum, Heidelberg,

69120, Germany

SOURCE:

AUTHOR(S):

Virology (1998), 240(2), 359-365 CODEN: VIRLAX; ISSN: 0042-6822

PUBLISHER: Academic Press

DOCUMENT TYPE:

Journal

English LANGUAGE:

The DNA genomes of four new human papillomaviruses, HPV 75, HPV 76, HPV 77, and HPV 80, have been cloned, sequenced, and characterized. HPV 75, HPV 76 (both HPV 49-related), and HPV 77 (HPV 29-related) were isolated from benign cutaneous warts and HPV 80 (HPV 15-related) from histol. normal skin. HPV 77 has also been

> 571-272-2528 Searcher : Shears

demonstrated in dysplastic warts and squamous cell carcinomas of the skin. The sequence data presented in this study led to a proposed modification of the definition of a new HPV type. The high degree of DNA sequence similarity between the E7 ORF of HPV 77 and HPV 29 (97.7%), as opposed to the E6 (82.8%) and L1 (85.3%) ORFs, might suggest conservation of a specific function or a possible recombinational event. Only the E6 and L1 ORFs of HPV 75 and HPV 76 have a similarity lower than 90%, whereas the DNA sequences of their upstream regulatory regions (URRs) share a similarity of 93%. The E7, E1, and E4 ORFs, as well as the URR of HPV 15 and HPV 80, share sequence similarities higher than 90%. Such a divergence in the similarity between different segments of the virus genomes of closely related HPV types has not been noted to date. A detailed comparative sequence anal. was performed. HPV 75, HPV 76, and HPV 80 revealed features characteristic of truly cutaneous HPV types, whereas HPV 77 shared several characteristics with the mucosal HPV types, some of which may have functional consequences.

IT 204338-84-1

RL: PRP (Properties)

(amino acid sequence; genomes of three of four novel HPV types, defined by differences of their L1 genes, show high conservation of E7 gene and URR)

REFERENCE COUNT:

THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 17 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

38

ED Entered STN: 05 Jun 1996

ACCESSION NUMBER: 1996:326231 HCAPLUS

DOCUMENT NUMBER: 125:8475

TITLE: Papilloma virus-like particles containing fusion

proteins for vaccines

INVENTOR(S): Gissmann, Lutz; Zhou, Jian; Mueller, Martin

PATENT ASSIGNEE(S): Medigene Gesellschaft fuer Molekularbiologische

Diagnostik, Therapie und Technologie Mbh, USA

SOURCE: Ger. Offen., 4 pp.

CODEN: GWXXBX

Patent

DOCUMENT TYPE:

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
DE 4435907	A1 19960411	DE 1994-4435907	19941007
DE 4435907	C2 19970724		
DE 4447664	C2 19990415	DE 1994-4447664	19941007
CA 2202090	AA 19960418	CA 1995-2202090	19951009
WO 9611272	A2 19960418	WO 1995-EP3974	19951009
WO 9611272	A3 19960926		
W: AU, BR, C	A, JP, MX, US		
RW: AT, BE, CI	H, DE, DK, ES, FR, GB	GR, IE, IT, LU,	MC, NL, PT,
SE			
AU 9642701	A1 19960502	AU 1996-42701	19951009
EP 809700	Al 19971203	EP 1995-934663	19951009
R: AT. BE. CI	H. DE. DK. ES. FR. GB	B, GR, IT, LI, LU,	NL, SE, MC,

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PT, IE
      JP 11504801
                        T2 19990511
                                           JP 1995-512335
                                                             19951009
                                           US 1997-817335
                                                             19971002
      US 6066324
                        Α
                             20000523
                             20020326
                                           US 1999-397680 19990916
      US 6361778
                        В1
                                            AU 2000-10106
                                                             20000105
      AU 760615
                        B2
                             20030522
      AU 2000010106
                        A5
                             20000316
                                           US 2001-949404
                             20030729
                                                             20010906
      US 6599508
                        В1
                                         DE 1994-4435907 A3 19941007
 PRIORITY APPLN. INFO.:
                                         DE 1995-19526752 A 19950721
                                         AU 1996-42701 A3 19951009
                                                        W 19951009
                                         WO 1995-EP3974
                                                        A3 19971002
A3 19990916
                                         US 1997-817335
                                         US 1999-397680
      Papilloma virus-like particles are prepared containing recombinant viral
 AΒ
      structural proteins L1 and/or L2 in which segments of these proteins
      are deleted and may be replaced with other proteins (e.g. viral
      early proteins) or protein fragments. These particles may be used
      in vaccines against tumors of the reproductive tract, e.g. cervical
      carcinoma.
. IT
      176521-52-1P
      RL: BAC (Biological activity or effector, except adverse); BPN
      (Biosynthetic preparation); BSU (Biological study, unclassified);
      THU (Therapeutic use); BIOL (Biological study); PREP (Preparation);
      USES (Uses)
         (papilloma virus-like particles containing fusion proteins
         for vaccines)
      ANSWER 18 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN
      Entered STN: 12 Jan 1996
 ACCESSION NUMBER:
                          1996:27856 HCAPLUS
 DOCUMENT NUMBER:
                          124:108357
 TITLE:
                          Sequence variation in the capsid protein genes
                          of human papillomavirus type 16 and type 31
                          Icenogle, Joseph P.; Clancy, Kelly A.; Lin,
 AUTHOR(S):
```

Sophia Y.

Public Health Service, U.S. Dep. of Health and CORPORATE SOURCE:

Human Services, Atlanta, GA, 30333, USA

Virology (1995), 214(2), 664-9 SOURCE:

CODEN: VIRLAX; ISSN: 0042-6822

Academic PUBLISHER: DOCUMENT TYPE: Journal LANGUAGE: English

The sequences of the capsid genes of a human papillomavirus type 16 (HPV 16) DNA and an HPV 31 DNA were determined The HPV 16 DNA contained genes coding for the most variable HPV 16 capsid proteins yet identified (17 variable amino acids). Three of six coding changes in the HPV 31 DNA occurred at positions equivalent to ones where variable amino acids in HPV 16 have been observed Variable amino acids in both viruses occurred predominantly in regions which showed amino acid variation when closely related types of HPV were compared; thus, most of the factors which determined the intratypic variation in the capsid proteins of the viruses described here were likely the same as those which determined differences between the capsid proteins of different HPV types.

IT 172892-77-2 172892-79-4 RL: PRP (Properties)

> Shears 571-272-2528 Searcher :

(amino acid sequence; sequence variation in capsid protein genes of human papillomavirus type 16 and type 31)

L4 ANSWER 19 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 02 Sep 1995

ACCESSION NUMBER: 1995:772980 HCAPLUS

DOCUMENT NUMBER: 123:225926

TITLE: Self-assembling recombinant papillomavirus

capsid proteins as vaccine and for diagnosis Lowy, Douglas R.; Schiller, John T.; Kirnbauer,

ADDITIONATION NO

DAME

INVENTOR(S): Lowy, Down Reinhard

PATENT ASSIGNEE(S): United States Dept. of Health and Human

Services, USA

SOURCE: U.S., 20 pp. Cont.-in-part of U.S. Ser. No. 941,

371.

WIND DAME

CODEN: USXXAM

DOCUMENT TYPE:

Patent English

LANGUAGE: En En FAMILY ACC. NUM. COUNT: 4

PATENT INFORMATION:

PA'	PATENT NO.					KIND DATE			A.	DATE						
US	5437951			A		1995	0801		U	s 19	93-3	19930316				
WO	9405	05792			l	1994	0317		W	o 19	93-U	S834	2	1993	0903	
	W:			JP												
	RW:	AT, SE	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IE,	IT,	LU	MC,	NL,	PT,
EP	6621			A:	1	1995	0712		E:	P 19	93-9	2135	3	1993	0903	
	R:	AT,	•			DK,										NL,
		PT,				1006	0503		-	- 10	00 5	0740	1	1002	0003	
	0850			T		1996					93-5			1993		
	6832			B		1997			A	0 19	93-4	84/5		1993	0903	
	9348					1994							-	1000	0000	
	2001		88			2001					01-1			1993		
	5618			A		1997					94-3			1994		
	5709			Α		1998					95-4	. —		1995		
	5716			A		1998					95-4			1995		
	5744			Α		1998					95-4			1995		
	5756			A		1998					95-4			1995		
	5871			A		1999					95-4		-	_		
	5985			Α		1999					95-4			1995		
	5855			A		1999					97-7			1997		
+	9944			Α.		1999			A	0 T 9	99-4	44/9		1999	0813	
	7179			B		2000				~ ^^			_	0001	0400	
	2003			A.		2003					01-8			2001		
	2002			A.		2002										
	2003			A.		2003			•		03-3		-	2003		
	2003			A.	L	2003	0911				03-4			2003		
PRIORIT	Y APP	LN.	INFO	. :							9413			1992		
											3286		A	1993		
											5074			1993		
•											US83			1993		
											3194			1994		
											4845			1995		
									AU 1	995-	3828	4	АЗ	1995	T006	

US 1997-781084 A1 19970109 US 1998-170129 B1 19981012 US 1999-316487 B1 19990521 US 2001-832065 B1 20010409 US 2001-878840 B1 20010611

AB Recombinant papillomavirus capsid proteins that are capable of self-assembly into capsomer structures and viral capsids that comprise conformational antigenic epitopes are provided. The capsomer structures and viral capsids, consisting of the capsid proteins that are expression products of a bovine, monkey or human papillomavirus L1 conformational coding sequence proteins, can be prepared as vaccines to induce a high-titer neutralizing antibody response in vertebrate animals. The self assembling capsid proteins can also be used as elements of diagnostic immunoassay procedures for papillomavirus infection. In example, described were construction and expression of recombinant papillomavirus L1 or L1/L2 capsid protein-encoding DNA, production of antisera in rabbits, purification of the bovine papillomavirus BPV1, BPV1 neutralization assay, and serum neutralizing titer against BPV1.

IT 155578-70-4

RL: PRP (Properties)

(amino acid sequence; self-assembling recombinant papillomavirus capsid proteins as vaccine and for diagnosis)

L4 ANSWER 20 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 03 Feb 1993

ACCESSION NUMBER: 1993:37211 HCAPLUS

DOCUMENT NUMBER: 118:37211

OCCUMENT NUMBER: 110.57211

TITLE: Induction of cytotoxic T lymphocytes with

peptides in vitro: Identification of candidate

T-cell epitopes in human papilloma virus Strauss, Hans J.; Davies, Huw; Sadovnikova,

Elena; Chain, Benny; Horowitz, Neil; Sinclair,

Christine

CORPORATE SOURCE:

SOURCE:

AUTHOR(S):

Imp. Cancer Res. Fund, Univ. Coll., London, UK Proceedings of the National Academy of Sciences

of the United States of America (1992), 89(17),

7871-5

CODEN: PNASA6; ISSN: 0027-8424

DOCUMENT TYPE:

Journal

LANGUAGE:

English

AB A set of overlapping peptides corresponding to the L1, E6, and E7 proteins of human papilloma virus 16 was tested for their ability to bind to major histocompatibility complex class I mols. and to stimulate cytotoxic T-lymphocyte (CTL) responses in vitro. A class I binding assay using intact RMA-S cells showed that 20 of the 99 human papilloma virus peptides bound to H-2Kb and/or Db mols. Fifteen of the 20 class I-binding peptides stimulated primary CTL responses, whereas peptides that were neg. in the binding assay failed to do so. Peptide-induced CTLs recognized the immunizing peptide very efficiently, requiring no more than 1-10 nM peptide for target cell lysis. However, 2 observations were made that have important implications for the design of peptide-based vaccines for inducing CTLs. Not all major histocompatibility complex-binding peptides that contained known motifs characteristic of naturally

processed peptides induced CTLs. The efficiency of CTL lysis was strongly decreased when the size of the target peptide differed by only 1 amino acid residue from that of the immunizing peptide. Thus, peptides chosen for vaccination must correspond in length to naturally processed peptides.

IT 143743-51-5

SOURCE:

RL: BIOL (Biological study)

(of L1 protein of human papilloma virus 16, structure of, in class I histocompatibility antigen binding and cytotoxic T-lymphocyte activation)

L4 ANSWER 21 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 01 Nov 1992

ACCESSION NUMBER: 1992:569102 HCAPLUS

DOCUMENT NUMBER: 117:169102

TITLE: Definition of linear antigenic regions of the

HPV16 L1 capsid protein using synthetic

virion-like particles

AUTHOR(S): Zhou, Jian; Sun, Xiao-Yi; Davies, Huw; Crawford,

Lionel; Park, David; Frazer, Ian H.

CORPORATE SOURCE: Lions Hum. Immunol. Lab., Univ. Queensland,

Woolloongabba, 4102, Australia Virology (1992), 189(2), 592-9

CODEN: VIRLAX; ISSN: 0042-6822

DOCUMENT TYPE: Journal LANGUAGE: English

Mice of 3 haplotypes (H-2d, H-2b, and H-2d/b) were immunized with synthetic human papillomavirus (HPV)16-like particles (VLPs), produced using a vaccinia virus doubly recombinant for the L1 and L2 proteins of HPV16. The resultant anti-VLP antisera recognized HPV16 capsids by ELISA assay and baculovirus recombinant HPV16 L1 and L2 protein on immunoblot. Overlapping peptides corresponding to the HPV16 L1 amino acid sequences were used to define the immunoreactive regions of the L1 protein. The majority of the L1 peptides were reactive with IgG from the mice immunized with the synthetic HPV16 capsids. A computer algorithm predicted 7 B epitopes in HPV16 L1, 5 of which lay within peptides strongly reactive with the murine antisera. The murine anti-VLP antisera failed to react with the 2 peptides recognized by anti-HPV16L1 monoclonal antibodies raised by others against recombinant L1 fusion protein. Thus, immunoreactive epitopes of HPV16 defined using virus-like particles differ significantly from those defined using recombinant HPV16 L1 fusion proteins, which implies that such fusion proteins may not be the antigens to look for in HPV16L1-specific immune responses in HPV-infected patients.

IT 143743-51-5P

RL: PREP (Preparation)

(of L1 capsid protein of human papillomavirus 16, preparation and antigenicity of)

L4 ANSWER 22 OF 22 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 31 Mar 1990

ACCESSION NUMBER: 1990:116820 HCAPLUS

DOCUMENT NUMBER: 112:116820

TITLE: Identification of immunogenic regions of the major coat protein of human papillomavirus type

16 that contain type-restricted epitopes Cason, John; Patel, Daksha; Naylor, Jennifer; AUTHOR(S): Lunney, Declan; Shepherd, Philip S.; Best, Jennifer M.; McCance, Dennis J. Richard Dimbleby Lab. Cancer Virology, London, CORPORATE SOURCE: SE1 7EH, UK Journal of General Virology (1989), 70(11), SOURCE: 2973-87 CODEN: JGVIAY; ISSN: 0022-1317 DOCUMENT TYPE: Journal English LANGUAGE: Regions were identified of the major capsid protein, L1, of the human papillomavirus (HPV) type 16 (HPV-16 L1), that are recognized by 5 monoclonal antibodies (MAbs) raised to a bacterial fusion protein containing residues 172-375 of HPV-16 L1. All 5 MAbs recognized HPV-16-infected tissue sections by immunohistochem., but not sections infected with HPV-la (cutaneous warts), HPV-6b or -11 (genital warts). MAbs 3D1, 5A4, and 1D6 also recognized HPV-2-infected sections (cutaneous warts); MAb 8C4 recognized only sections containing HPV-16. Four MAbs (8C4, 3D1, 1D6, and 5A4) recognized a synthetic peptide corresponding to residues 269-284 of HPV-16 L1; within this region a min. antibody binding site was identified, a tripeptide 276-278. However, the complete epitope appears to extend beyond these residues and beyond HPV-16 L1 (269-284). The 5th MAb, 1C6, recognized bacterial fusion proteins containing HPV-6b L1, -16 L1 or -18 L1 using immunoblots, yet appeared HPV-16-specific when tested on infected tissue sections. This MAb recognized 5 amino acids within a different region of HPV-16 L1 (residues 299-313). IT 125551-62-4 RL: BIOL (Biological study) (from human papilloma virus type 16, preparation and antigenic determinant mapping on) E30 THROUGH E59 ASSIGNED FILE 'REGISTRY' ENTERED AT 15:30:11 ON 03 MAR 2004 30 SEA FILE=REGISTRY ABB=ON PLU=ON (143743-51-5/BI OR L5 243135-53-7/BI OR 308800-09-1/BI OR 125551-62-4/BI OR 155578-70-4/BI OR 172892-77-2/BI OR 172892-79-4/BI OR 176521-52-1/BI OR 204338-84-1/BI OR 244773-43-1/BI OR 272100-89-7/BI OR 272100-91-1/BI OR 295372-67-7/BI OR 308800-07-9/BI OR 344806-30-0/BI OR 405194-67-4/BI OR 434341-62-5/BI OR 434341-76-1/BI OR 475437-53-7/BI OR 475437-55-9/BI OR 582340-39-4/BI OR 583085-22-7/BI OR 622878-55-1/BI OR 624440-70-6/BI OR 627561-74-4/BI OR 627561-77-7/BI OR 627561-78-8/BI OR 627561-79-9/BI OR 627561-80-2/BI OR 627561-81-3/BI) 30 L1 AND L5 L6 ANSWER 1 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN L6 627561-81-3 REGISTRY RN Protein (synthetic human papillomavirus 16 clone H) (9CI) (CA INDEX NAME) OTHER NAMES:

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571-272-2528

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Searcher :

Shears 571-272-2528

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          L-Lysine, L-seryl-L-alanyl-L-tyrosyl-L-alanyl-L-alanyl-L-asparaginyl-
CN
          L-alanylglycyl-L-valyl-L-\alpha-aspartyl-L-asparaginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-arginyl-L-ar
          α-glutamyl-L-cysteinyl-L-isoleucyl-L-seryl-L-methionyl-L-
          α-aspartyl-L-tyrosyl- (9CI) (CA INDEX NAME)
OTHER NAMES:
          13: PN: WO0244384 SEQID: 13 unclaimed sequence
CN
SQL
         20
SEO
                   1 SAYAANAGVD NRECISMDYK
```

==== =====

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7-15
HITS AT:
REFERENCE
          1: 137:19374
    ANSWER 15 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
1.6
     405194-67-4 REGISTRY
RN
    Antigen VCA (viral capsid antigen) (human papillomavirus 16 strain
CN
     114/B gene L1h) (9CI) (CA INDEX NAME)
OTHER NAMES:
    Capsid protein (synthetic HPV16 strain 114/B gene 16L1h)
CN
    Protein (human papillomavirus 16 strain 114/B gene L1p)
CN
    Protein (synthetic HPV16 strain 114/B gene 16L1p)
CN
CI
SOL
    505
         1 MSLWLPSEAT VYLPPVPVSK VVSTDEYVAR TNIYYHAGTS RLLAVGHPYF
SEO
        51 PIKKPNNNKI LVPKVSGLQY RVFRIHLPDP NKFGFPDTSF YNPDTQRLVW
       101 ACVGVEVGRG QPLGVGISGH PLLNKLDDTE NASAYAANAG VDNRECISMD
       151 YKQTQLCLIG CKPPIGEHWG KGSPCTNVAV NPGDCPPLEL INTVIQDGDM
       201 VDTGFGAMDF TTLQANKSEV PLDICTSICK YPDYIKMVSE PYGDSLFFYL
       251 RREQMFVRHL FNRAGAVGEN VPDDLYIKGS GSTANLASSN YFPTPSGSMV
       301 TSDAQIFNKP YWLQRAQGHN NGICWGNQLF VTVVDTTRST NMSLCAAIST
       351 SETTYKNTNF KEYLRHGEEY DLQFIFQLCK ITLTADVMTY IHSMNSTILE
       401 DWNFGLQPPP GGTLEDTYRF VTSQAIACQK HTPPAPKEDP LKKYTFWEVN
       451 LKEKFSADLD QFPLGRKFLL QAGLKAKPKF TLGKRKATPT TSSTSTTAKR
       501 KKRKL
          139-147, 304-312
HITS AT:
**RELATED SEQUENCES AVAILABLE WITH SEQLINK**
REFERENCE
            1: 136:257935
    ANSWER 16 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
L6
     344806-30-0 REGISTRY
RN
     Protein L1 (human papillomavirus 31) (9CI) (CA INDEX NAME)
OTHER NAMES:
     46: PN: WO0141799 PAGE: 23-24 claimed sequence
CN
CI
    MAN
SQL 504
         1 MSLWRPSEAT VYLPPVPVSK VVSTDEYVTR TNIYYHAGSA RLLTVGHPYY
SEO
        51 SIPKSDNPKK IWVPKVSGLQ YRVFRVRLPD PNKFGFPDTS FYNPETQRLV
       101 WACVGLEVGR GQPLGVGISG HPLLNKFDDT ENSNRYAGGP GTDNRECISM
       151 DYKOTOLCLL GCKPPIGEHW GKGSPCSNNA ITPGDCPPLE LKNSVIQDGD
       201 MVDTGFGAMD FTALQDTKSN VPLDICNSIC KYPDYLKMVA EPYGDTLFFY
       251 LRREQMFVRH FFNRSGTVGE SVPTDLYIKG SGSTATLANS TYFPTPSGSM
       301 VTSDAQIFNK PYWMQRAQGH NNGICWGNQL FVTVVDTTRS TNMSVCAAIA
       351 NSDTTFKSSN FKEYLRHGEE FDLQFIFQLC KITLSADIMT YIHSMNPAIL
       401 EDWNFGLTTP PSGSLEDTYR FVTSQAITCQ KTAPQKPKED PFKDYVFWEV
       451 NLKEKFSADL DQFPLGRKFL LQAGYRARPK FKAGKRSAPS ASTTTPAKRK
       501 KTKK
HITS AT:
          305-313
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Shears

Searcher :

571-272-2528

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1: 135:45179
REFERENCE
    ANSWER 17 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
L6
    308800-09-1 REGISTRY
RN
    L-Isoleucine, L-alanylglycyl-L-valyl-L-\alpha-aspartyl-L-
    asparaginyl-L-arginyl-L-\alpha-glutamyl-L-cysteinyl- \  \, (9CI)
                                                             (CA
    INDEX NAME)
OTHER NAMES:
   2: PN: DE19925235 SEQID: 2 claimed protein
SQL 9
         1 AGVDNRECI
SEQ
           =======
HITS AT:
           1-9
REFERENCE
          1: 138:383747
            2: 134:16537
REFERENCE
    ANSWER 18 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
1.6
    308800-07-9 REGISTRY
RN
    L-Tryptophan, L-alanyl-L-glutaminyl-L-isoleucyl-L-phenylalanyl-L-
    asparaginyl-L-lysyl-L-prolyl-L-tyrosyl- (9CI) (CA INDEX NAME)
OTHER NAMES:
    1: PN: DE19925235 SEQID: 1 claimed protein
CN
SQL
         1 AQIFNKPYW
SEO
HITS AT:
           1-9
REFERENCE
           1: 134:16537
    ANSWER 19 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
L6
    295372-67-7 REGISTRY
RN
    Protein L1 (human papillomavirus 16) (9CI) (CA INDEX NAME)
CN
OTHER NAMES:
    1: PN: WO0054730 SEQID: 1 claimed protein
CI
    MAN
SQL 531
         1 MOVTFIYILV ITCYENDVNV YHIFFQMSLW LPSEATVYLP PVPVSKVVST
SEO
        51 DEYVARTNIY YHAGTSRLLA VGHPYFPIKK PNNNKILVPK VSGLQYRVFR
       101 IHLPDPNKFG FPDTSFYNPD TQRLVWACVG VEVGRGQPLG VGISGHPLLN
       151 KLDDTENASA YAANAGVDNR ECISMDYKQT QLCLIGCKPP IGEHWGKGSP
                          ===========
       201 CTNVAVNPGD CPPLELINTV IQDGDMVHTG FGAMDFTTLQ ANKSEVPLDI
       251 CTSICKYPDY IKMVSEPYGD SLFFYLRREQ MFVRHLFNRA GTVGENVPDD
       301 LYIKGSGSTA NLASSNYFPT PSGSMVTSDA QIFNKPYWLQ RAQGHNNGIC
       351 WGNQLFVTVV DTTRSTNMSL CAAISTSETT YKNTNFKEYL RHGEEYDLQF
       401 IFQLCKITLT ADVMTYIHSM NSTILEDWNF GLQPPPGGTL EDTYRFVTQA
       451 IACOKHTPPA PKEDDPLKKY TFWEVNLKEK FSADLDQFPL GRKFLLQAGL
       501 KAKPKFTLGK RKATPTTSST STTAKRKKRK L
HITS AT:
           165-173, 330-338
```

RELATED SEQUENCES AVAILABLE WITH SEQLINK REFERENCE 1: 133:256735 ANSWER 20 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN L6 272100-91-1 REGISTRY RN Protein (synthetic papillomavirus biotin-binding protein 169) (9CI) CN (CA INDEX NAME) OTHER NAMES: 5: PN: WO0031128 SEQID: 7 claimed protein CN CI 499 SOL 1 MSLWLPSEAT VYLPPVPVSK VVSTDEYVAR TNIYYHAGTS RLLAVGHPYF SEO 51 PIKKPNNNKI LVPKVSGLQY RVFRIHLPDP NKFGFPDTSF YNPDTQRLVW 101 ACVGVEVGRG QPLGVGISGH PLLNKLDDTE NASAYAANAG VDNRECISMD 151 YKQTQLCLIG CKPPIGEHWG KGSPCTNVAV NPGDCPPLEL INTVIQDGDM 201 VDTGFGAMDF TTLQANKSEV PLDICTSICK YPDYIKMVSE PYGDSLFFYL 251 RREQMFVRHL FNRAGAVGEN VPDDLYIKGS GSTANLASSN YFPTPSGSMV 301 TSDAQIFNKP YWLQRAQGHN NGICWGNQLF VTVVDTTRST NMSLCAAIST 351 SETTYKNTNF KEYLRHGEEY DLQFIFQLCK ITLTADVMTY IHSMNSTILE 401 DWNFGLQPPP GGTLEDTYRF VTSQAIASQK HTPPAPKEDP LKKYTFWEVN 451 LKEKFSADLD QFPLGRKFLL QAGLKAKPKF TLGGGGRGEF TGTYITAVT HITS AT: 139-147, 304-312 1: 133:22397 REFERENCE ANSWER 21 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN L6 272100-89-7 REGISTRY RN Protein (synthetic papillomavirus biotin-binding protein 168) (9CI) CN (CA INDEX NAME) OTHER NAMES: 3: PN: WO0031128 SEQID: 5 claimed protein CN CI MAN SQL 497 1 MSLWLPSEAT VYLPPVPVSK VVSTDEYVAR TNIYYHAGTS RLLAVGHPYF SEO 51 PIKKPNNNKI LVPKVSGLQY RVFRIHLPDP NKFGFPDTSF YNPDTQRLVW 101 ACVGVEVGRG OPLGVGISGH PLLNKLDDTE NASAYAANAG VDNRECISMD 151 YKQTQLCLIG CKPPIGEHWG KGSPCTNVAV NPGDCPPLEL INTVIQDGDM 201 VDTGFGAMDF TTLQANKSEV PLDICTSICK YPDYIKMVSE PYGDSLFFYL 251 RREQMFVRHL FNRAGAVGEN VPDDLYIKGS GSTANLASSN YFPTPSGSMV 301 TSDAQIFNKP YWLQRAQGHN NGICWGNQLF VTVVDTTRST NMSLCAAIST ====== == 351 SETTYKNTNF KEYLRHGEEY DLQFIFQLCK ITLTADVMTY IHSMNSTILE 401 DWNFGLQPPP GGTLEDTYRF VTSQAIASQK HTPPAPKEDP LKKYTFWEVN 451 LKEKFSADLD QFPLGRKFLL QAGLKAKPKF TLGGGGCSWA PPFKASC 139-147, 304-312 HITS AT: 1: 133:22397 REFERENCE ANSWER 22 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN 1.6

Searcher :

571-272-2528

Shears

```
244773-43-1 REGISTRY
RN
     1-473-Protein L 1 (human papillomavirus 16 gene L1) fusion protein
CN
    with 1-55-RNA formation factor (human papillomavirus 16 gene E7)
     (9CI) (CA INDEX NAME)
OTHER NAMES:
    PN: WO9948917 PAGE: 12 claimed sequence
CI
    MAN
SOL
    528
         1 MSLWLPSEAT VYLPPVPVSK VVSTDEYVAR TNIYYHAGTS RLLAVGHPYF
SEO
        51 PIKKPNNNKI LVPKVSGLQY RVFRIHLPDP NKFGFPDTSF YNPDTQRLVW
       101 ACVGVEVGRG QPLGVGISGH PLLNKLDDTE NASAYAANAG VDNRECISMD
       151 YKQTQLCLIG CKPPIGEHWG KGSPCTNVAV NPGDCPPLEL INTVIQDGDM
       201 VHTGFGAMDF TTLQANKSEV PLDICTSICK YPDYIKMVSE PYGDSLFFYL
       251 RREOMFVRHL FNRAGTVGEN VPDDLYIKGS GSTANLASSN YFPTPSGSMV
       301 TSDAQIFNKP YWLQRAQGHN NGICWGNQLF VTVVDTTRST NMSLCAAIST
       351 SETTYKNTNF KEYLRHGEEY DLQFIFQLCK ITLTADVMTY IHSMNSTILE
       401 DWNFGLOPPP GGTLEDTYRF VTSQAIACQK HTPPAPKEDP LKKYTFWEVN
       451 LKEKFSADLD QFPLGRKFLL QAGMHGDTPT LHEYMLDLQP ETTDLYCYEQ
       501 LNDSSEEEDE IDGPAGQAEP DRAHYNIV
          139-147, 304-312
HITS AT:
            1: 131:262610
REFERENCE
    ANSWER 23 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
L6
     243135-53-7 REGISTRY
RN
     Capsid protein (human papillomavirus 16 SiHa cell variant) (9CI)
CN
     (CA INDEX NAME)
OTHER NAMES:
CN
     32: PN: WO0141799 PAGE: 21 claimed sequence
CN
    Protein L1 (human papillomavirus 16)
CI
    MAN
SOL 531
         1 MQVTFIYILV ITCYENDVNV YHIFFQMSLW LPSEATVYLP PVPVSKVVST
SEQ
        51 DEYVARTNIY YHAGTSRLLA VGHPYFPIKK PNNNKILVPK VSGLQYRVFR
       101 IHLPDPNKFG FPDTSFYNPD TQRLVWACVG VEVGRGQPLG VGISGHPLLN
       151 KLDDTENASA YAANAGVDNR ECISMDYKQT QLCLIGCKPP IGEHWGKGSP
                          ==========
       201 CTNVAVNPGD CPPLELINTV IQDGDMVDTG FGAMDFTTLQ ANKSEVPLDI
       251 CTSICKYPDY IKMVSEPYGD SLFFYLRREQ MFVRHLFNRA GAVGENVPDD
       301 LYIKGSGSTA NLASSNYFPT PSGSMVTSDA QIFNKPYWLQ RAQGHNNGIC
                                          = =======
       351 WGNQLFVTVV DTTRSTNMSL CAAISTSETT YKNTNFKEYL RHGEEYDLQF
       401 IFQLCKITLT ADVMTYIHSM NSTILEDWNF GLQPPPGGTL EDTYRFVTSQ
       451 AIACQKHTPP APKEDPLKKY TFWEVNLKEK FSADLDQFPL GRKFLLQAGL
       501 KAKPKFTLGK RKATPTTSST STTAKRKKRK L
          165-173, 330-338
HITS AT:
**RELATED SEQUENCES AVAILABLE WITH SEQLINK**
            1: 135:45179
REFERENCE
REFERENCE
          2: 131:209883
```

Searcher :

Shears

571-272-2528

```
L6
    ANSWER 24 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
RN
     204338-84-1 REGISTRY
     Protein (human papillomavirus 77 gene L1) (9CI) (CA INDEX NAME)
CN
OTHER NAMES:
CN
     GenBank CAA75468
     GenBank CAA75468 (Translated from: GenBank Y15175)
CN
CI
SQL
    565
         1 MCIYTLAPTL FCLLLHNGLL FLYYLLTOHI MCTLMEAIFI CGLLPFLCLG
SEO
        51 NVAVNVFHIF LQMALWRSSD NLVYLPPTPV SKVISTDDYV TRTNVYYYAG
       101 SSRLLTVGHP YFAIPKTSGT KVDVPKVSAF QYRVFRVRLP DPNKFGLPDA
       151 RIYNPEAERL VWACTGVEVG RGQPLGVGLS GHPLYNKLND TENSNIAHAD
       201 NSPDSRDNIS VDCKQTQLCI LGCTPPMGEY WGKGTPCART NTTPGDCPPL
       251 ELMTSYIQDG DMVDTGYGAM DFTALQFNKS DVPLDICQSI CKYPDYLGMA
       301 ADPYGDSMFF FLRREQLFAR HFFNRAGDVG DKIPESLYLK GSSGRETPGS
       351 AIYSPTPSGS MVTSEAQIFN KPYWLQQAQG HNNGICWGNQ VFLTVVDTTR
                           ===== ====
       401 STNMSLSAST ESQTPSTYDA TKIKEYLRHG EEYDLQFIFQ LCKVTLTPEI
       451 MAYIHTMNTA LLEDWNFGLT LPPSTSLEDT YRFVTSSAIT CQKDVAPTEK
       501 QDPYAKLNEW DVDLKDRETL DLSQFPLGRK FLLQIGARRR SVVPSRKRRA
       551 PTPSPASTKR KRSKK
          366-374
HITS AT:
          1: 128:213916
REFERENCE
     ANSWER 25 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
L6
RN
     176521-52-1 REGISTRY
     (1-305)-(316-505)-Protein L1 (human papillomavirus 16 coat) (9CI)
CN
     (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN
     (1-305)-(316-505)-Protein L1 (human papilloma virus 16 coat)
CI
SQL 495
         1 MSLWLPSEAT VYLPPVPVSK VVSTDEYVAR TNIYYHAGTS RLLAVGHPYF
SEQ
        51 PIKKPNNNKI LVPKVSGLQY RVFRIYLPDP NKFGFPDTSF YNPDTQRLVW
       101 ACVGVEVGRG OPLGVGISGH PLLNKLDDTE NASAYAANAG VDNRECISMD
       151 YKOTOLCLIG CKPPIGEHWG KGSPCNNVAV TPGDCPPLEL INTVIQDGDM
       201 VDTGFGAMDF TTLQANKSEV PLDICTSICK YPDYIKMVSE PYGDSLFFYL
       251 RREQMFVRHL FNRAGAVGEN VPDDLYIKGS GPTANLASSN YFPTPSGSMV
       301 TSDAQAQGHN NGICWGNQLF VTVVDTTRST NMSLCAAIST SEPTYKNTNF
       351 KEYLRHGEEY DLQFIFQLCK ITLTADVMTY IHSMNSTILE DWNFGLQPPP
       401 GGTLEDTYRF VTSQAIACQK HTPPAPKEDP LKKYTFWEVN LKEKFSADLD
       451 QFPLGRKFLL QAGFKAKPKF TLGKRKATPT TSSTSTTAKR KKRKL
HITS AT:
          139-147
REFERENCE
            1: 125:8475
     ANSWER 26 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
L6
     172892-79-4 REGISTRY
RN
     Protein L1 (human papillomavirus 31 capsid) (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     Protein L1 (human papilloma virus 31 capsid)
```

Searcher :

Shears

571-272-2528

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OTHER NAMES:
    Capsid protein L1 (human papillomavirus 31)
    L1 capsid protein (human papillomavirus 31)
CN
CI
    MAN
SQL 504
         1 MSLWRPSEAT VYLPPVPVSK VVSTDEYVTR TNIYYHAGSA RLLTVGHPYY
SEO
        51 SIPKSDNPKK IVVPKVSGLQ YRVFRVRLPD PNKFGFPDTS FYNPETQRLV
       101 WACVGLEVGR GOPLGVGISG HPLLNKFDDT ENSNRYAGGP GTDNRECISM
       151 DYKOTOLCLL GCKPPIGEHW GKGSPCSNNA ITPGDCPPLE LKNSVIQDGD
       201 MVDTGFGAMD FTALQDTKSN VPLDICNSIC KYPDYLKMVA EPYGDTLFFY
       251 LRREOMFVRH FFNRSGAVGE SVPNDLYIKG SGSTATLANS TYFPTPSGSM
       301 VTSDAOIFNK PYWMQRAQGH NNGICWGNQL FVTVVDTTRS TNMSVCAAIA
               ____
       351 NSDTTFKSSN FKEYLRHGEE FDLQFIFQLC KITLSADIMT YIHSMNPAIL
       401 EDWNFGLTTP PSGSLEDTYR FVTSQAITCQ KTAPQKPKED PFKDYVFWEV
       451 NLKEKFSADL DQFPLGRKFL LQAGYRARPK FKAGKRSAPS ASTTTPAKRK
       501 KTKK
HITS AT:
          305-313
**RELATED SEQUENCES AVAILABLE WITH SEQLINK**
REFERENCE
           1: 124:108357
    ANSWER 27 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
L6
     172892-77-2 REGISTRY
RN
    Protein L1 (human papillomavirus 16 capsid) (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
    Protein L1 (human papilloma virus 16 capsid)
CN
OTHER NAMES:
    Capsid protein L1 (human papillomavirus 16)
CN
CN
    L1 coat protein (human papillomavirus 16)
CI
SOL 505
        1 MSLWLPSEAT VYLPPVPVSK VVSTDEYVAR TNIYYHAGTS RLLAVGHPYF
SEO
        51 PIKKPNNNKI LVPKVSGLQY RVFRIYLPDP NKFGFPDTSF YNPDTQRLVW
       101 ACVGVEVGRG OPLGVGISGH PLLNKLDDTE NASAYAANAG VDNRECISMD
       151 YKQTQLCLIG CKPPIGEHWG KGSPCNNVAV TPGDCPPLEL INTVIQDGDM
       201 VDTGFGAMDF TTLQANKSEV PLDICTSICK YPDYIKMVSE PYGDSLFFYL
       251 RREQMFVRHL FNRAGAVGEN VPDDLYIKGS GPTANLASSN YFPTPSGSMV
       301 TSDAQIFNKP YWLQRAQGHN NGICWGNQLF VTVVDTTRST NMSLCAAIST
              ======= ==
       351 SEPTYKNTNF KEYLRHGEEY DLQFIFQLCK ITLTADVMTY IHSMNSTILE
       401 DWNFGLOPPP GGTLEDTYRF VTSQAIACQK HTPPAPKEDP LKKYTFWEVN
       451 LKEKFSADLD QFPLGRKFLL QAGFKAKPKF TLGKRKATPT TSSTSTTAKR
       501 KKRKL
          139-147, 304-312
HITS AT:
**RELATED SEQUENCES AVAILABLE WITH SEQLINK**
REFERENCE
            1: 124:108357
    ANSWER 28 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
L6
     155578-70-4 REGISTRY
RN
```

Searcher :

Shears 571-272-2528

```
Protein L 1 (bovine papillomavirus 16 gene L1 reduced) (9CI)
                                                                                                                                                                                                                                         (CA
CN
                 INDEX NAME)
OTHER CA INDEX NAMES:
                Protein L 1 (bovine papilloma virus 16 gene L1 reduced)
OTHER NAMES:
                 Protein (human papilloma 16 virus strain wild-type gene L1)
CN
                 MAN
CT
SQL
                505
                               1 MSLWLPSEAT VYLPPVPVSK VVSTDEYVAR TNIYYHAGTS RLLAVGHPYF
SEQ
                            51 PIKKPNNNKI LVPKVSGLQY RVFRIHLPDP NKFGFPDTSF YNPDTQRLVW
                        101 ACVGVEVGRG QPLGVGISGH PLLNKLDDTE NASAYAANAG VDNRECISMD
                        151 YKOTOLCLIG CKPPIGEHWG KGSPCTNVAV NPGDCPPLEL INTVIQDGDM
                        201 VDTGFGAMDF TTLQANKSEV PLDICTSICK YPDYIKMVSE PYGDSLFFYL
                        251 RREQMFVRHL FNRAGTVGEN VPDDLYIKGS GSTANLASSN YFPTPSGSMV
                        301 TSDAQIFNKP YWLQRAQGHN NGICWGNQLF VTVVDTTRST NMSLCAAIST
                        351 SETTYKNTNF KEYLRHGEEY DLQFIFQLCK ITLTADVMTY IHSMNSTILE
                        401 DWNFGLQPPP GGTLEDTYRF VTQAIACQKH TPPAPKEDDP LKKYTFWEVN
                        451 LKEKFSADLD QFPLGRKFLL QAGLKAKPKF TLGKRKATPT TSSTSTTAKR
                        501 KKRKL
                                      139-147, 304-312
HITS AT:
REFERENCE
                                         1:
                                                    123:225926
REFERENCE
                                         2:
                                                       121:5012
                 ANSWER 29 OF 30 REGISTRY COPYRIGHT 2004 ACS on STN
1.6
                 143743-51-5 REGISTRY
RN
                 L-Methionine, L-tyrosyl-L-alanyl-L-alanyl-L-asparaginyl-L-
CN
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